

CLAIMS

Subt 5

Subt 10

Subt 15

Subt 20

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Subt 95

1. A method of assaying for an analyte which method comprises the steps of:

- i) mixing a sample of cells possibly containing the analyte with a cell lysis reagent to provide a cell lysis fluid,
- ii) mixing the cell lysis fluid with reagents, including a specific binding partner of the analyte for binding to the analyte, for performing a specific binding assay for the analyte,
- iii) and mixing the cell lysis fluid with a sequestrant for the cell lysis reagent, whereby the binding of step ii) is performed in the presence of the sequestrant.

15 2. A method as claimed in claim 1, wherein the cell lysis reagent is a detergent.

3. A method as claimed in claim 1, wherein the sequestrant is a cyclodextrin.

4. A method as claimed in claim 3, wherein the amount of

20 sequestrant is in the range of 1 - 5% of the binding reaction mixture.

5. A method as claimed in claim 1, wherein steps i), ii) and iii) are all performed in a single reaction vessel.

6. A method as claimed in claim 1, wherein multiple assays are performed in parallel in wells of a multiwell plate.

25 7. A method as claimed in claim 1, wherein the cells are cultured in a vessel and are lysed in that vessel for assaying the analyte in that vessel.

8. A method as claimed in claim 1, wherein the assay of step ii) is a homogenous assay.

30 9. A method as claimed in claim 1, wherein the assay of step ii) is a scintillation proximity assay.

10. A method as claimed in claim 1, wherein the specific binding assay of step ii) is an immunoassay.

11. A method as claimed in claim 1, wherein the analyte is adenosine-3',5'-cyclic monophosphate, the cell lysis reagent is dodecyl trimethyl ammonium bromide and the sequestrant is α -cyclodextrin.

5 12. A method as claimed in claim 1, wherein the cells have been maintained in a culture medium, and step i) is performed in the presence of the culture medium.

13. A method as claimed in claim 1, wherein the intracellular or 10 the total (intracellular plus extracellular) concentration is measured of an analyte selected from adenosine-3',5'-cyclic monophosphate, interleukin-6 and prostaglandin E₂.

14. A kit, suitable for assaying for an analyte by the method as 15 claimed in claim 1, comprising: a detergent; a sequestrant for the detergent; a specific binding partner of the analyte; a tracer; and separation means for separating bound tracer from unbound tracer.

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